

REMARKS

Claims 1-6 were pending in the application and were rejected. Claims 1-6 are herein amended.

Preliminary Comments

The Amendment filed on December 19, 2007 included a Declaration by the inventor. The experimental data referred to in the Declaration was inadvertently omitted. Thus, attached hereto is a re-submission of the Amendment and Declaration, with the experimental data included.

Article 19 Amendments

From the statement in the last paragraph of page 3 of the Office Action "...the limitation 'for a non-viral gene delivery sector' is functional language ... ", it appears that the U.S.P.T.O. is not examining the claims as amended in the Article 19 Amendments. The U.S.P.T.O. electronic file wrapper has an entry labeled "claims" which contains the original claims and the Article 19 amended claims.

In view of the above, Applicant respectfully requests that the U.S.P.T.O. consider the Article 19 amended claims. For clarity, Applicant herewith submits a clean copy of the claims including the Article 19 amendments.

Objections to the Claims

Claims 1-4 are objected to as allegedly containing "informalities."

The Office Action states that in the description of R₇ semicolons are used to separate the members of the group rather than commas. In response, Applicant herein amends claims 1-4 to correct for this and other informalities. Favorable reconsideration is respectfully

requested.

Applicant's Response to Claim Rejections under 35 U.S.C. §102

Claims 1 and 2 were rejected under 35 U.S.C. 102(b) as allegedly being anticipated by Onishi (US Pat. 4,816,540).

It is the position of the Office Action that that Onishi discloses Applicant's claimed copolymers and that the functional language "for a non-viral gene delivery sector" does not materially change the structure of the polymer.

The Office Action considers the cationic graft-copolymers of Onishi to fully encompass Applicant's claims 1 and 2. That is, the Office Action's position is that the cationic graft-copolymers of Onishi meet the recited cationic graft-copolymers and that such cationic graft-copolymers also inherently would be capable of the claimed function of being a non-viral gene delivery vector.

In response, Applicant herein amends claims 1 and 2 in order to recite "a non-viral gene delivery vector formed of an aqueous solution of a cationic graft-copolymer..." Onishi does not disclose or suggest such an aqueous solution. Favorable reconsideration is respectfully requested.

Applicant's Response to Claim Rejections under 35 U.S.C. §103

Claims 3-6 were rejected under 35 U.S.C. 103(a) as allegedly being unpatentable over Onishi (US Pat. 4,816,540) in view of Pack, Gene-Delivery Polymers.

It is the position of the Office Action that Onishi discloses the invention as claimed, wit the exception of teaching forming a complex with DNA. The Office Action cites Pack

as teaching forming a complex between a cationic graft polymer and DNA (Section 2.2.2).

Onishi claims a cationic graft-copolymer, useful as a micro carrier for cell cultivation, comprising a unit derived from a cationic polysaccharide whereas the present invention claims a cationic graft-copolymer of a water-soluble linear backbone polymer having hydroxyl groups comprising a unit derived from a cationic a water-soluble linear polysaccharide. Although both the inventions are directed at a cationic graft-copolymer, Onishi does not disclose or teach or suggest any possibility for a non-viral gene delivery vector by the graft copolymer. Also, a person having ordinary skill in the art would not have been prompted to make use of a cationic water-soluble linear polysaccharide in the preparation of cationic graft-copolymer.

Though the Office Action cites Pack (Gene-delivery polymers) as teaching forming a complex between a cationic graft polymer and DNA (Section 2.2.2), Pack also shows that the polymers are not copolymers, but rather homopolymers by the characterization that the polymers themselves comprise linear, branched, and dendrimeric structures. On the other hand, a cationic graft-copolymer of this invention is characterized as copolymer having a hydrophilic-hydrophobic micro-separated-domain as shown in the specification.

In section 2.2 Pack also shows various synthetic vector, including DEAE-dextran of a starting material for this invention, has suffered from problems including toxicity, low gene transfer efficiency, and *in vivo* instability.

It would not have been obvious that a cationic graft-copolymer obtained by graft-polymerizing methyl methacrylate of an olefin monomer onto DEAE-dextran of a cationic derivative of a water-soluble linear polymer having hydroxyl groups can solve these problems, especially of toxicity and low gene transfer efficiency. Samples 1 having a 150%

weight increase, 2 having a 200% weight increase and 3 having a 300% weight increase were prepared following procedure of Example 1.

With the transfection efficiency, transfection activity was determined using the X-gal staining (β -galactosidase activities in tissue) method and a value 3 times higher was confirmed for sample 1 and sample 2 than for the starting DEAE-dextran hydrochloride.

Formation of a complex between nucleic acids (DNA or RNA) and cationic graft-copolymers, such as DEAE-dextran-MMA copolymer, is improved as compared with DEAE-dextran of the starting material.

In these Examples, a complex of sample 1 between DNA and DEAE-dextran-MMA-Copolymer hydrochloride having a 150% weight increase was formed in 2 hours. A complex of sample 2 and sample 3 between DNA and DEAE-Dextran-MMA-copolymer hydrochloride having a 200% and 300% weight increase were formed in 1 hour and 0.5 hours, respectively. However, a complex between DNA and DEAE-dextran hydrochloride was formed in 96 hours.

Additionally, Applicant herewith submits experimental data which shows that the efficacy/transfection efficiency of the claimed non-viral gene delivery vector comprising of the graft copolymer compared with a prior known vector. Accordingly, Applicant respectfully submits that the attached data is sufficient to prove the higher efficacy/efficiency of a transfection with the graft copolymer by comparison of a starting material.

Specifically, Applicant submits the data of Example 1(DDMC) compared to PolyFect Reagent (QIAGEN GmbH) having dendrimeric structures shown in Section 2.2.2 of Pack (Gene-delivery polymers). The data is for the transfection for COS7 cells by

pGL3DNA/DDMC. The cells were ready to harvest 72 hours after transfection, and were assayed for luciferase activity.

Accordingly, in view of the above remarks and the cited art, Applicant respectfully submits that the claimed gene delivery of polymers would not have been obvious to one having ordinary skill in the art. Favorable reconsideration is respectfully requested.

Applicant's Response to Double Patenting Rejections

Claims 1 and 2 were rejected on the ground of nonstatutory obviousness-type double patenting as allegedly being unpatentable over claim 1 of U.S. Patent No. 4,816,540.

Claims 3-6 were rejected on the ground of nonstatutory obviousness-type double patenting as allegedly being unpatentable over claim 1 of U.S. Patent No. 4,816,540 in view of Pack, Gene-Delivery Polymers.

In response, Applicant respectfully submits that since the §102 and §103 rejections are overcome on the technical merits, the obviousness-type double patenting rejections are also overcome. Favorable reconsideration is respectfully requested.

Conclusion

In view of the aforementioned amendments and accompanying remarks, Applicant submits that the claims, as herein amended, are in condition for allowance. Applicant requests such action at an early date.

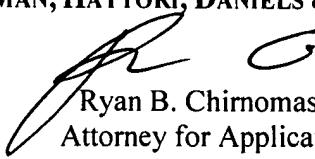
If the Examiner believes that this application is not now in condition for allowance, the Examiner is requested to contact Applicant's undersigned attorney to arrange for an

interview to expedite the disposition of this case.

If this paper is not timely filed, Applicant respectfully petitions for an appropriate extension of time. The fees for such an extension or any other fees that may be due with respect to this paper may be charged to Deposit Account No. 50-2866.

Respectfully submitted,

WESTERMAN, HATTORI, DANIELS & ADRIAN, LLP



Ryan B. Chirnomas
Attorney for Applicant
Registration No. 56,527
Telephone: (202) 822-1100
Facsimile: (202) 822-1111

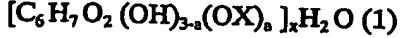
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Enclosure: Clean copy of Article 19 Amendments corresponding to claims 1-4
Declaration under 37 CFR 1.132 (including experimental data)

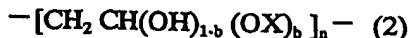
CLAIMS

What is claimed is:

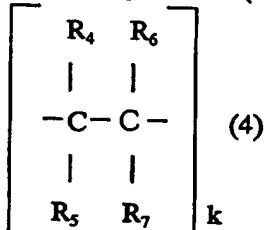
1. A cationic graft-copolymer of a water-soluble linear backbone polymer having hydroxyl groups, for a non-viral gene delivery vector, comprising a unit derived from a cationic water-soluble linear polysaccharide of the following formula (1)



or a unit derived from a water-soluble linear polyvinylalcohol of the following formula (2) or a partial hydrolyzed alcohol of the following formula (3)



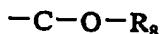
Wherein X is a $-(CH_2)_mR_1$ organic radical where R_1 is a member of the class consisting of $-NH_2$ radical, $-N(CH_3)_2$ radical, $-N(C_2H_5)_2$ radical, $-N^+(C_2H_5)_3$ radical, $-N^+(CH_2)_2CH_2CH(OH)CH_3$ radical, $-N^+(C_2H_5)_2CH_2CH(OH)CH_3$ radical, $-N^+(C_2H_5)_2(C_2H_5)N(C_2H_5)_2$ radical, $-C_6H_4NH_2$ radical, and $-COC_6H_4NH_2$ radical, $-COR_2$ radical where R_2 is $-CH_2NH_2$ or $-C_6H_4NH_2$, $-CH_2CH(OH)CH_2R_3$ radical where R_3 is $-NH_2$, $-N(CH_3)_2$, $-N(C_2H_5)_2$, and $-N^+(C_2H_5)_3$ radical, m is a natural number of 1 to 3, a is a positive number having a value of $0 < a < 3$, b is a positive number having a value of $0 < b < 1$, x and n are natural numbers having a value of 5 or more, $1 > b + c$, and Ac is acetyl radical; and a unit derived from a polymerize-able olefin compound of the following formula (4)



Wherein R_4 , R_5 and R_6 are each selected from the group consisting of hydrogen and CH_3 and R_7 is a member of the group consisting of

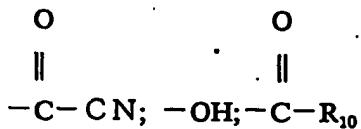
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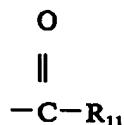


Where R_8 is a member of the class consisting of hydrogen, C_1-C_{12} alkyl radicals, cyclohexyl radical, C_1-C_4 hydroxyalkyl radicals, C_1-C_8 aminoalkyl radicals, C_1-C_8 dialkylaminoalkyl radicals, glycidyl radical, tetrahydrofuran radical, C_1-C_4 lower alkyl-substituted tetrahydrofuran radical, benzyl radical, the $(CH_2CH_2O)_yCH_2CH_2OH$ radical where y is a positive integer from 1 to 10, and $-N(R_9)_2$ where the two R_9 's which may be the same or different, are

either hydrogen or a $C_1 - C_4$ alkyl radical;

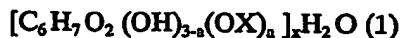


Where R_{10} is a $C_1 - C_8$ alkyl radical; phenyl radical; tolyl radical; pyridine radical; pyrrolidone radical; and

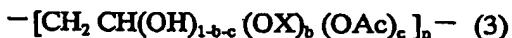
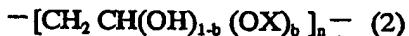


Where R_{11} is NH_2 , NHCH_3 , N,N-dimethylamino radical, $\text{N,N-dimethylaminopropylamino}$ radical, and morpholine radical.

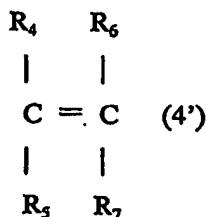
2. A process for preparing a cationic graft-copolymer of a water-soluble linear backbone polymer having hydroxyl groups, for a non-viral gene delivery vector, which comprises reacting a cationic water-soluble linear polysaccharide of the following formula (1)



or a unit derived from a water-soluble linear polyvinylalcohol of the following formula (2) or a partial hydrolyzed alcohol of the following formula (3)

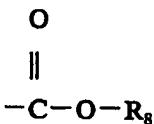


Wherein X is a $-(\text{CH}_2)_m\text{R}_1$ organic radical where R_1 is a member of the class consisting of $-\text{NH}_2$ radical, $-\text{N}(\text{CH}_3)_2$ radical, $-\text{N}(\text{C}_2\text{H}_5)_2$ radical, $-\text{N}^+(\text{C}_2\text{H}_5)_3$ radical, $-\text{N}^+(\text{CH}_2)_2\text{CH}_2\text{CH}(\text{OH})\text{CH}_3$ radical, $-\text{N}^+(\text{C}_2\text{H}_5)_2\text{CH}_2\text{CH}(\text{OH})\text{CH}_3$ radical, $-\text{N}^+(\text{C}_2\text{H}_5)_2(\text{C}_2\text{H}_5)\text{N}(\text{C}_2\text{H}_5)_2$ radical, $-\text{C}_6\text{H}_4\text{NH}_2$ radical, and $-\text{COC}_6\text{H}_4\text{NH}_2$ radical, $-\text{COR}_2$ radical where R_2 is $-\text{CH}_2\text{NH}_2$ or $-\text{C}_6\text{H}_4\text{NH}_2$, $-\text{CH}_2\text{CH}(\text{OH})\text{CH}_2\text{R}_3$ radical where R_3 is $-\text{NH}_2$, $-\text{N}(\text{CH}_3)_2$, $-\text{N}(\text{C}_2\text{H}_5)_2$, and $-\text{N}^+(\text{C}_2\text{H}_5)_3$ radical, m is a natural number of 1 to 3, a is a positive number having a value of $0 < a < 3$, b is a positive number having a value of $0 < b < 1$, x and n are natural numbers having a value of 5 or more, $1 > b+c$, and Ac is acetyl radical; with a polymerize-able olefin compound of the formula (4')

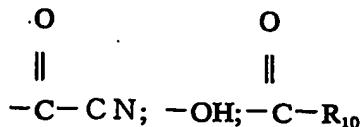


Wherein R_4 , R_5 , and R_6 are each selected from the group consisting of hydrogen and CH_3

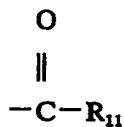
and R₇ is a member of the group consisting of



Where R₈ is a member of the class consisting of hydrogen, C₁ — C₁₂ alkyl radicals, cyclohexyl radical, C₁ — C₄ hydroxyalkyl radicals, C₁ — C₈ aminoalkyl radicals, C₁ — C₈ dialkylaminoalkyl radicals, glycidyl radical, tetrahydrofuran radical, C₁ — C₄ lower alkyl -substituted tetrahydrofuran radical, benzyl radical, the (CH₂CH₂ O)_y CH₂CH₂OH radical where y is a positive integer from 1 to 10, and —N(R₉)₂ where the two R₉s which may be the same or different, are either hydrogen or a C₁ — C₄ alkyl radical;



Where R₁₀ is a C₁ — C₈ alkyl radical; phenyl radical; tolyl radical; pyridine radical; pyrrolidone radical; and

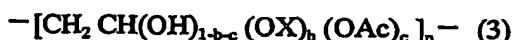
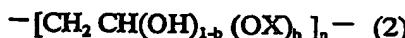


Where R₁₁ is NH₂, NHCH₃, N,N-dimethylamino radical, N,N-dimethylaminopropylamino radical, and morpholine radical.

3. A complex between a cationic graft-copolymer of a water-soluble linear backbone polymer having hydroxyl groups and DNA, comprising a unit derived from a cationic water-soluble linear polysaccharide of the following formula (1)



or a unit derived from a water-soluble linear polyvinylalcohol of the following formula (2) or a partial hydrolyzed alcohol of the following formula (3)



Wherein X is a —(CH₂)_m R₁ organic radical where R₁ is a member of the class consisting of

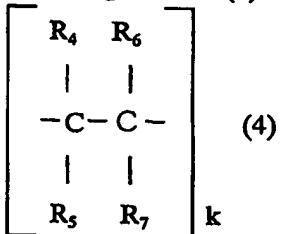
—NH₃⁺ radical, —NH⁺(CH₃)₂ radical, —NH⁺(C₂H₅)₂ radical, —N⁺(C₂H₅)₃ radical,

—N⁺(CH₂)₂CH₂CH(OH)CH₃ radical, —N⁺(C₂H₅)₂CH₂CH(OH)CH₃ radical,

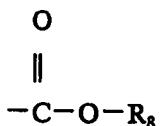
—N⁺(C₂H₅)₂(C₂H₅)N(C₂H₅)₂ radical, —C₆H₄NH₃⁺ radical, and —COC₆H₄NH₃⁺ radical,

—COR₂ radical where R₂ is —CH₂NH₃⁺ or —C₆H₄NH₃⁺, —CH₂CH(OH)CH₂R₃ radical

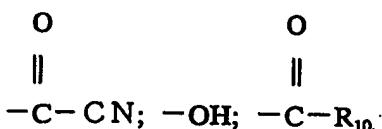
where R_3 is $-NH_3^+$, $-NH^+(CH_3)_2$, $-NH^+(C_2H_5)_2$, and $-N^+(C_2H_5)_3$ radical, m is a natural number of 1 to 3, a is a positive number having a value of $0 < a < 3$, b is a positive number having a value of $0 < b < 1$, x and n are natural numbers having a value of 5 or more, $1 > b + c$, and Ac is acetyl radical; a unit derived from a polymerizable olefin compound of the following formula (4)



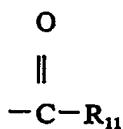
Wherein R_4 , R_5 and R_6 are each selected from the group consisting of hydrogen and CH_3 , and R_7 is a member of the group consisting of



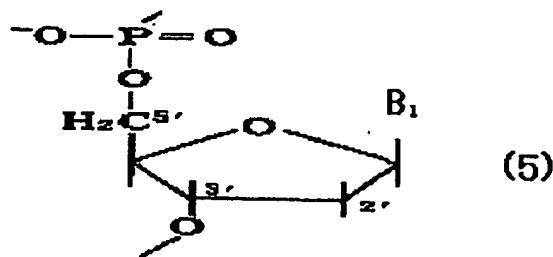
Where R_8 is a member of the class consisting of hydrogen, $C_1 - C_{12}$ alkyl radicals, cyclohexyl radical, $C_1 - C_4$ hydroxyalkyl radicals, $C_1 - C_8$ aminoalkyl radicals, $C_1 - C_8$ dialkylaminoalkyl radicals, glycidyl radical, tetrahydrofuran radical, $C_1 - C_4$ lower alkyl-substituted tetrahydrofuran radical, benzyl radical, the $(CH_2CH_2O)_y$, CH_2CH_2OH radical where y is a positive integer from 1 to 10, and $-N(R_9)_2$ where the two R_9 's which may be the same or different, are either hydrogen or a $C_1 - C_4$ alkyl radical;



Where R_{10} is a $C_1 - C_8$ alkyl radical; phenyl radical; tolyl radical; pyridine radical; pyrrolidone radical; and

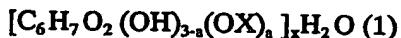


Where R_{11} is NH_2 , $NHCH_3$, N,N -dimethylamino radical, N,N -dimethylaminopropylamino radical, and morpholine radical; and a unit derived from a poly(deoxyribonucleotide) of the following formula (5) as a recurring unit.

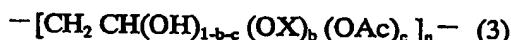
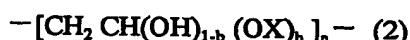


Where B_1 is a base selected from the group of adenine, thymine, guanine, and cytosine.

4. A complex between a cationic graft-copolymer of a water-soluble linear backbone polymer having hydroxyl groups and RNA, comprising a unit derived from a cationic water-soluble linear polysaccharide of the following formula (1)



or a unit derived from a water-soluble linear polyvinylalcohol of the following formula (2) or a partial hydrolyzed alcohol of the following formula (3)



Wherein X is a $-(CH_2)_mR_1$ organic radical where R_1 is a member of the class consisting of

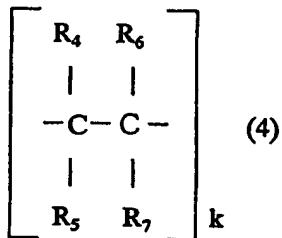
$-NH_3^+$ radical, $-NH^+(CH_3)_2$ radical, $-NH^+(C_2H_5)_2$ radical, $-N^+(C_2H_5)_3$ radical,

$-N^+(CH_2)_2CH_2CH(OH)CH_3$ radical, $-N^+(C_2H_5)_2CH_2CH(OH)CH_3$ radical,

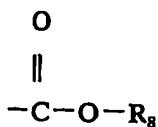
$-N^+(C_2H_5)_2(C_2H_5)N(C_2H_5)_2$ radical, $-C_6H_4NH_3^+$ radical, and $-COC_6H_4NH_3^+$ radical,

$-COR_2$ radical where R_2 is $-CH_2NH_3^+$ or $-C_6H_4NH_3^+$, $-CH_2CH(OH)CH_2R_3$ radical

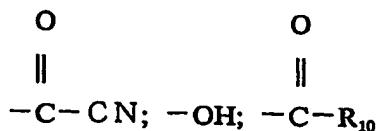
where R_3 is $-NH_3^+$, $-NH^+(CH_3)_2$, $-NH^+(C_2H_5)_2$, and $-N^+(C_2H_5)_3$ radical, m is a natural number of 1 to 3, a is a positive number having a value of $0 < a < 3$, b is a positive number having a value of $0 < b < 1$, x and n are natural numbers having a value of 5 or more, $1 > b+c$, and Ac is acetyl radical; a unit derived from a polymerizable olefin compound of the following formula (4)



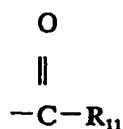
Wherein R_4 , R_5 and R_6 are each selected from the group consisting of hydrogen and CH_3 and R_7 is a member of the group consisting of



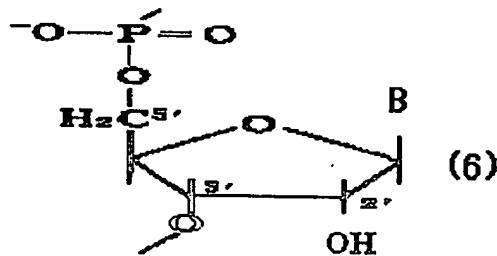
Where R_8 is a member of the class consisting of hydrogen, C_1-C_{12} alkyl radicals, cyclohexyl radical; C_1-C_4 hydroxyalkyl radicals, C_1-C_8 aminoalkyl radicals, C_1-C_8 dialkylaminoalkyl radicals, glycidyl radical, tetrahydrofuran radical, C_1-C_4 lower alkyl -substituted tetrahydrofuran radical, benzyl radical, the $(\text{CH}_2\text{CH}_2\text{O})_y\text{CH}_2\text{CH}_2\text{OH}$ radical where y is a positive integer from 1 to 10, and $-\text{N}(\text{R}_9)_2$ where the two R_9 s which may be the same or different, are either hydrogen or a C_1-C_4 alkyl radical;



Where R_{10} is a C_1-C_8 alkyl radical; phenyl radical; tolyl radical; pyridine radical; pyrrolidone radical; and



Where R_{11} is NH_2 , NHCH_3 , N,N-dimethylamino radical, $\text{N,N-dimethylaminopropylamino}$ radical, and morpholine radical; and a unit derived from a poly(ribonucleotide) of the following formula(6) as a recurring unit.



Where B is a base selected from the group of adenine, uracil, guanine, and cytosine.

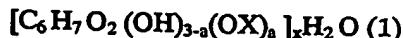
5. A gene delivery system using a complex between the cationic graft-copolymer and DNA, of Claim 3.
6. A gene delivery system using a complex between the cationic graft-copolymer and RNA, of Claim 4.

AMENDED CLAIMS

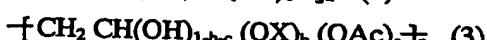
[Received by the International Bureau on 01 October 2004 (01.10.04):
original claims 1-4 are amended and all other claims are retained unchanged. (6 pages)]

What is claimed is:

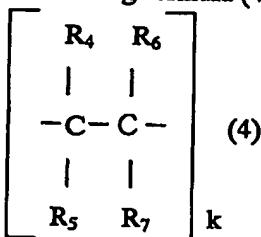
1. (amended) A non-viral gene delivery vector formed from a cationic graft-copolymer of a water-soluble linear backbone polymer having hydroxyl groups comprising a unit derived from a cationic water-soluble linear polysaccharide of the following formula (1)



or a unit derived from a water-soluble linear polyvinylalcohol of the following formula (2) or a partial hydrolyzed alcohol of the following formula (3)



Wherein X is a $-(CH_2)_m R_1$ organic radical where R_1 is a member of the class consisting of $-NH_2$ radical, $-N(CH_3)_2$ radical, $-N(C_2H_5)_2$ radical, $-N^+(C_2H_5)_3$ radical, $-N^+(CH_2)_2CH_2CH(OH)CH_3$ radical, $-N^+(C_2H_5)_2CH_2CH(OH)CH_3$ radical, $-N^+(C_2H_5)_2(C_2H_5)N(C_2H_5)_2$ radical, $-C_6H_4NH_2$ radical, and $-CO-C_6H_4NH_2$ radical, $-COR_2$ radical where R_2 is $-CH_2NH_2$ or $-C_6H_4NH_2$, $-CH_2CH(OH)CH_2R_3$ radical where R_3 is $-NH_2$, $-N(CH_3)_2$, $-N(C_2H_5)_2$, and $-N^+(C_2H_5)_3$ radical, m is a natural number of 1 to 3, a is a positive number having a value of $0 < a < 3$, b is a positive number having a value of $0 < b < 1$, x and n are natural numbers having a value of 5 or more, $1 > b + c$, and Ac is acetyl radical; and a unit derived from a polymerize-able olefin compound of the following formula (4)



Wherein R_4 , R_5 and R_6 are each selected from the group consisting of hydrogen and CH_3 and R_7 is a member of the group consisting of

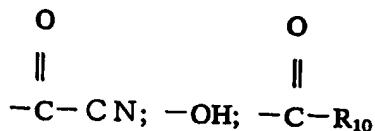
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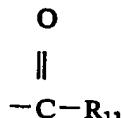


Where R_8 is a member of the class consisting of hydrogen, C_1-C_{12} alkyl radicals, cyclohexyl radical, C_1-C_4 hydroxyalkyl radicals, C_1-C_8 aminoalkyl radicals, C_1-C_8 dialkylaminoalkyl radicals, glycidyl radical, tetrahydrofuran radical, C_1-C_4 lower alkyl-substituted tetrahydrofuran radical, benzyl radical, the $(CH_2CH_2O)_yCH_2CH_2OH$ radical where y is a positive integer from 1 to 10, and $-N(R_9)_2$ where the two R_9 's which may be the same or different, are

either hydrogen or a C₁—C₄ alkyl radical;



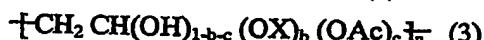
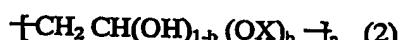
Where R₁₀ is a C₁—C₈ alkyl radical; phenyl radical; tolyl radical; pyridine radical; pyrrolidone radical; and



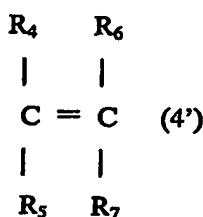
Where R₁₁ is NH₂, NHCH₃, N,N-dimethylamino radical, N,N-dimethylaminopropylamino radical, and morpholine radical.

2. (amended) A process for preparing a non-viral gene delivery vector formed from a cationic graft-copolymer of a water-soluble linear backbone polymer having hydroxyl groups which comprises reacting a cationic water-soluble linear polysaccharide of the following formula (1) [C₆H₇O₂(OH)_{3-a}(OX)_a]_xH₂O (1)

or a unit derived from a water-soluble linear polyvinylalcohol of the following formula (2) or a partial hydrolyzed alcohol of the following formula (3)

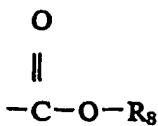


Wherein X is a $-(\text{CH}_2)_m\text{R}_1$ organic radical where R₁ is a member of the class consisting of —NH₂ radical, —N(CH₃)₂ radical, —N(C₂H₅)₂ radical, —N⁺(C₂H₅)₃ radical, —N⁺(CH₂)₂CH₂CH(OH)CH₃ radical, —N⁺(C₂H₅)₂CH₂CH(OH)CH₃ radical, —N⁺(C₂H₅)₂(C₂H₅)N(C₂H₅)₂ radical, —C₆H₄NH₂ radical, and —COC₆H₄NH₂ radical, —COR₂ radical where R₂ is —CH₂NH₂ or —C₆H₄NH₂, —CH₂CH(OH)CH₂R₃ radical where R₃ is —NH₂, —N(CH₃)₂, —N(C₂H₅)₂, and —N⁺(C₂H₅)₃ radical, m is a natural number of 1 to 3, a is a positive number having a value of 0<a<3, b is a positive number having a value of 0<b<1, x and n are natural numbers having a value of 5 or more, 1>b+c, and Ac is acetyl radical; with a polymerize-able olefin compound of the formula (4')

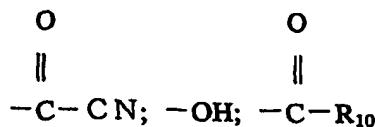


Wherein R₄, R₅ and R₆ are each selected from the group consisting of hydrogen and CH₃

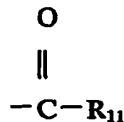
and R₇ is a member of the group consisting of



Where R₈ is a member of the class consisting of hydrogen, C₁—C₁₂ alkyl radicals, cyclohexyl radical, C₁—C₄ hydroxyalkyl radicals, C₁—C₈ aminoalkyl radicals, C₁—C₈ dialkylaminoalkyl radicals, glycidyl radical, tetrahydrofuran radical, C₁—C₄ lower alkyl-substituted tetrahydrofuran radical, benzyl radical, the (CH₂CH₂O)_yCH₂CH₂OH radical where y is a positive integer from 1 to 10, and—N(R₉)₂ where the two R₉s which may be the same or different, are either hydrogen or a C₁—C₄ alkyl radical;

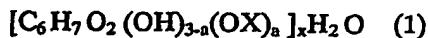


Where R₁₀ is a C₁—C₈ alkyl radical; phenyl radical; tolyl radical; pyridine radical; pyrrolidone radical; and

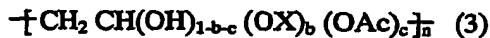


Where R₁₁ is NH₂, NHCH₃, N,N-dimethylamino radical, N,N-dimethylaminopropylamino radical, and morpholine radical.

3. (amended) A non-viral gene delivery vector, as the first step of transfection, using a complex between a cationic graft-copolymer of a water-soluble linear backbone polymer having hydroxyl groups and DNA, comprising a unit derived from a cationic water-soluble linear polysaccharide of the following formula (1)

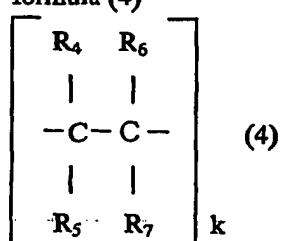


or a unit derived from a water-soluble linear polyvinylalcohol of the following formula (2) or a partial hydrolyzed alcohol of the following formula (3)

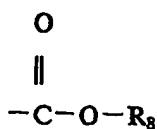


Wherein X is a —(CH₂)_n R₁ organic radical where R₁ is a member of the class consisting of —NH₃⁺ radical, —NH⁺ (CH₃)₂ radical, —NH⁺ (C₂H₅)₂ radical, —N⁺ (C₂H₅)₃ radical, —N⁺(CH₂)₂CH₂CH(OH)CH₃ radical, —N⁺(C₂H₅)₂CH₂CH(OH)CH₃ radical, —N⁺(C₂H₅)₂(C₂H₅)N(C₂H₅)₂ radical, —C₆H₄NH₃⁺ radical, and —COC₆H₄NH₃⁺ radical,

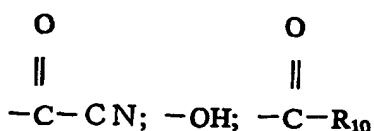
$-\text{COR}_2$ radical where R_2 is $-\text{CH}_2\text{NH}_3^+$ or $-\text{C}_6\text{H}_4\text{NH}_3^+$, $-\text{CH}_2\text{CH}(\text{OH})\text{CH}_2\text{R}_3$ radical where R_3 is $-\text{NH}_3^+$, $-\text{NH}^+(\text{CH}_3)_2$, $-\text{NH}^+(\text{C}_2\text{H}_5)_2$, and $-\text{N}^+(\text{C}_2\text{H}_5)_3$ radical, m is a natural number of 1 to 3, a is a positive number having a value of $0 < a < 3$, b is a positive number having a value of $0 < b < 1$, x and n are natural numbers having a value of 5 or more, $1 > b + c$, and Ac is acetyl radical; a unit derived from a polymerizable olefin compound of the following formula (4)



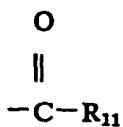
Wherein R_4 , R_5 and R_6 are each selected from the group consisting of hydrogen and CH_3 and R_7 is a member of the group consisting of



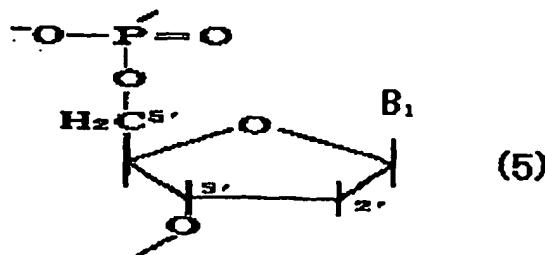
Where R_8 is a member of the class consisting of hydrogen, C_1-C_{12} alkyl radicals, cyclohexyl radical, C_1-C_4 hydroxyalkyl radicals, C_1-C_8 aminoalkyl radicals, C_1-C_8 dialkylaminoalkyl radicals, glycidyl radical, tetrahydrofuran radical, C_1-C_4 lower alkyl-substituted tetrahydrofuran radical, benzyl radical, the $(\text{CH}_2\text{CH}_2\text{O})_y\text{CH}_2\text{CH}_2\text{OH}$ radical where y is a positive integer from 1 to 10, and $-\text{N}(\text{R}_9)_2$ where the two R_9 's which may be the same or different, are either hydrogen or a C_1-C_4 alkyl radical;



Where R_{10} is a C_1-C_8 alkyl radical; phenyl radical; tolyl radical; pyridine radical; pyrrolidone radical; and

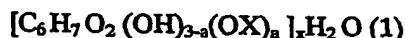


Where R_{11} is NH_2 , NHCH_3 , N,N-dimethylamino radical, $\text{N,N-dimethylaminopropylamino}$ radical, and morpholine radical; and a unit derived from a poly(deoxyribonucleotide) of the following formula (5) as a recurring unit.

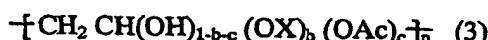
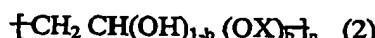


Where B_1 is a base selected from the group of adenine, thymine, guanine, and cytosine.

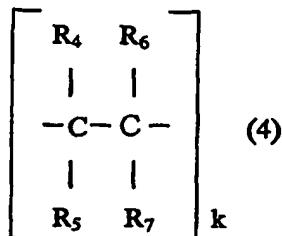
4. (amended) A non-viral gene delivery vector, as the first step of transfection, using a complex between a cationic graft-copolymer of a water-soluble linear backbone polymer having hydroxyl groups and RNA, comprising a unit derived from a cationic water-soluble linear polysaccharide of the following formula (1)



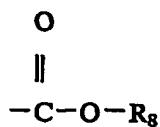
or a unit derived from a water-soluble linear polyvinylalcohol of the following formula (2) or a partial hydrolyzed alcohol of the following formula (3)



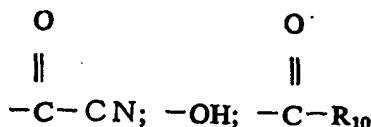
Wherein X is a $-(CH_2)_m R_1$ organic radical where R_1 is a member of the class consisting of $-NH_3^+$ radical, $-NH^+(CH_3)_2$ radical, $-NH^+(C_2H_5)_2$ radical, $-N^+(C_2H_5)_3$ radical, $-N^+(CH_2)_2CH_2CH(OH)CH_3$ radical, $-N^+(C_2H_5)_2CH_2CH(OH)CH_3$ radical, $-N^+(C_2H_5)_2(C_2H_5)N(C_2H_5)_2$ radical, $-C_6H_4NH_3^+$ radical, and $-COC_6H_4NH_3^+$ radical, $-COR_2$ radical where R_2 is $-CH_2NH_3^+$ or $-C_6H_4NH_3^+$, $-CH_2CH(OH)CH_2R_3$ radical where R_3 is $-NH_3^+$, $-NH^+(CH_3)_2$, $-NH^+(C_2H_5)_2$, and $-N^+(C_2H_5)_3$ radical, m is a natural number of 1 to 3, a is a positive number having a value of $0 < a < 3$, b is a positive number having a value of $0 < b < 1$, x and n are natural numbers having a value of 5 or more, $1 > b+c$, and Ac is acetyl radical; a unit derived from a polymerizable olefin compound of the following formula (4)



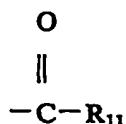
Wherein R_4 , R_5 and R_6 are each selected from the group consisting of hydrogen and CH_3 and R_7 is a member of the group consisting of



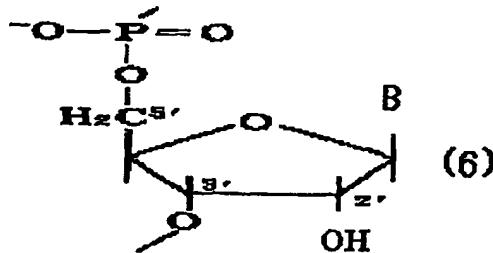
Where R_8 is a member of the class consisting of hydrogen, C_1-C_{12} alkyl radicals, cyclohexyl radical, C_1-C_4 hydroxyalkyl radicals, C_1-C_8 aminoalkyl radicals, C_1-C_8 dialkylaminoalkyl radicals, glycidyl radical, tetrahydrofuran radical, C_1-C_4 lower alkyl-substituted tetrahydrofuran radical, benzyl radical, the $(\text{CH}_2\text{CH}_2\text{O})_y\text{CH}_2\text{CH}_2\text{OH}$ radical where y is a positive integer from 1 to 10, and $-\text{N}(\text{R}_9)_2$ where the two R_9 's which may be the same or different, are either hydrogen or a C_1-C_4 alkyl radical;



Where R_{10} is a C_1-C_8 alkyl radical; phenyl radical; tolyl radical; pyridine radical; pyrrolidone radical; and



Where R_{11} is NH_2 , NHCH_3 , N,N-dimethylamino radical, $\text{N,N-dimethylaminopropylamino}$ radical, and morpholine radical; and a unit derived from a poly(ribonucleotide) of the following formula(6) as a recurring unit.



Where B is a base selected from the group of adenine, uracil, guanine, and cytosine.

Brief Statement

What is claimed by amendment for claim 1: Claim 1 is verified to be a non-viral gene delivery vector formed from a cationic graft-copolymer of formula(1) or formula(2) or formula(3) as detailed in application claim 1.

What is claimed by amendment for claim 2: Claim 2 is verified to be a process for preparing a non-viral gene delivery vector formed from a cationic graft-copolymer as described in application claim 2.

What is claimed by amendment for claim 3: Claim 3 is verified to be a non-viral gene delivery vector, as the first step of transfection, using a complex between DNA and a cationic graft-copolymer as described in application claim 3.

What is claimed by amendment for claim 4: Claim 4 is verified to be a non-viral gene delivery vector, as the first step of transfection, using a complex between RNA and a cationic graft-copolymer as described in application claim 4.